



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification: B01L 3/00	A1	(11) International Publication Number: (43) International Publication Date:	WO 00/25921 11 May 2000 (11.05.2000)
(21) International Application Number: PCT/SE99/01958			
(22) International Filing Date: 29 October 1999 (29.10.1999)		Published	
(30) Priority Data: 9803734-4 30 October 1998 (30.10.1998) SE			

(60) Parent Application or Grant

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(54) Title: LIQUID MICROVOLUME HANDLING SYSTEM

(54) Titre: SYSTEME DE TRAITEMENT D'UN MICROVOLUME DE LIQUIDE

(57) Abstract

The present invention relates to a microfluidic device comprising a microchannel (2, 4) providing for solvent contact between an open microarea (MA) carrying a microvolume (1) of a solvent and a reservoir (3) for the solvent, said reservoir (3) and said microchannel (2, 4) being adapted so that solvent evaporated from said microarea (MA) is continuously replaced by solvent from the reservoir (3) through said microchannel (2, 4). It further relates to method for replacing solvents evaporating from a microvolume (1) of solvent placed in an open microarea (MA) of a microfluidic device, wherein replacement is continuously taking place via a microchannel (2, 4) that transports solvent to the microarea (MA) from a solvent reservoir (vessel) (3). The device and method are suitable for preventing the desiccation of samples.

(57) Abrégé

L'invention porte sur un dispositif microfluidique comprenant un microcanal (2, 4) prévu pour que le solvant vienne en contact entre une microsurface (MA) ouverte supportant un microvolume (1) de solvant et un réservoir (3) de solvant, le réservoir (3) et le microcanal (2, 4) étant adaptés pour que le solvant s'évaporant de la microsurface (MA) soit remplacé en continu par le solvant provenant du réservoir (3) et passant par le microcanal (2, 4). L'invention porte également sur un procédé permettant de remplacer les solvants s'évaporant d'un microvolume (1) de solvant placé dans une microsurface (MA) ouverte d'un dispositif microfluidique, le remplacement s'effectuant en continu par un microcanal (2, 4) qui achemine le solvant vers la microsurface (MA) depuis le réservoir (réceptacle) (3) de solvant. Le dispositif et ce procédé sont également appropriés pour prévenir le dessèchement des échantillons.

PCT

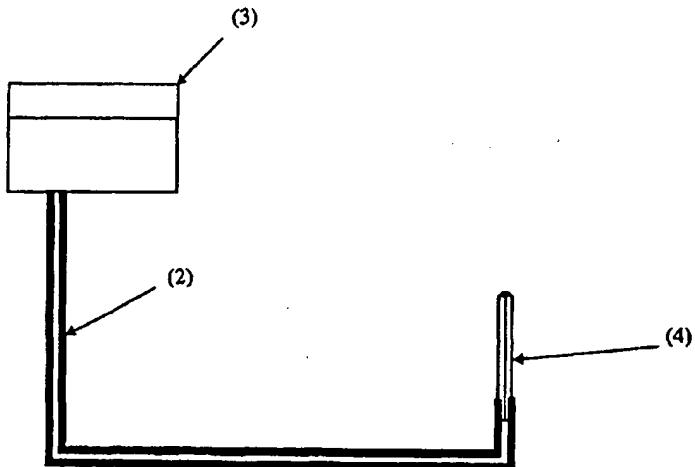
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(21) International Application Number: PCT/SE99/01958		(81) Designated States: JP, US, European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).
(22) International Filing Date: 29 October 1999 (29.10.99)		
(30) Priority Data: 9803734-4 30 October 1998 (30.10.98) SE		Published <i>With international search report.</i>
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(54) Title: LIQUID MICROVOLUME HANDLING SYSTEM



(57) Abstract

The present invention relates to a microfluidic device comprising a microchannel (2, 4) providing for solvent contact between an open microarea (MA) carrying a microvolume (1) of a solvent and a reservoir (3) for the solvent, said reservoir (3) and said microchannel (2, 4) being adapted so that solvent evaporated from said microarea (MA) is continuously replaced by solvent from the reservoir (3) through said microchannel (2, 4). It further relates to method for replacing solvents evaporating from a microvolume (1) of solvent placed in an open microarea (MA) of a microfluidic device, wherein replacement is continuously taking place via a microchannel (2, 4) that transports solvent to the microarea (MA) from a solvent reservoir (vessel) (3). The device and method are suitable for preventing the desiccation of samples.

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Description

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LIQUID MICROVOLUME HANDLING SYSTEM

TECHNICAL FIELD

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s The present invention relates to microfluidic devices comprising microchannels, and to methods for replacing solvent amounts that evaporate from open microareas carrying microvolumes of the solvent. The resent invention also relates to the use of the method for replacing solvent in a method for preventing the desiccation of samples. The microvolume of solvent may be in the form of a droplet (microdrop).

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BACKGROUND ART

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Microvolume handling systems have attracted a considerable interest in biochemical analysis, combinatorial chemistry and high throughput screening (HTS) applications. The miniaturised format is compatible in size with many interesting

issues of bioanalytical work, such as single cell analysis, when material is available only in extremely limited amounts. Furthermore, by decreasing the volume, an enhanced efficiency in terms of a higher rate of mixing and/or chemical reaction can be expected in the sample container, since the effect of diffusion and thermal

convection is more pronounced on a smaller scale.

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In HTS applications, goals are currently set on screening more than 10^5 compounds in a single assay. To manage such a tremendous number of samples with reasonable space, cost and time requirements, the miniaturised microtitre plate

format has been developed. Based on micromachining of different materials, e.g., by anisotropically etching single crystalline silicon wafers, well-defined picolitre to nanolitre vials are readily fabricated (Jansson et al. (1992) *J. Chromatography* 626, 310-314; Beyer Hietpas et al. (1995) *J. Liq. Chromatography* 18, 3557-3576).

Biomolecules such as DNA and proteins have been assayed in the microvial format utilising capillary electrophoresis (Jansson et al. *supra*; Beyer Hietpas et al., *supra*), bioluminescence (Croscheck et al. (1997) *Anal. Chem.* 69, 4768-4772), electrochemical analysis (Clark et al. (1997) *Anal. Chem.* 69, 259-263; Clark et al. (1998) *Anal. Chem.* 70, 1119-1125) and mass spectrometry (Jespersen et al. (1994) *J. Rapid Comm. in Mass Spectrom.* 8, 581-584).

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However, the rate of solvent evaporation is particularly pronounced for microvolumes, for instance small droplets, since the surface-to-volume ratio increases when the drop diameter decreases. The most common way for avoiding desiccation is by covering the containers with a material non-permeable for the underlying solvent. However, covers, either liquid or solid, inherently have the potential to introduce interfering compounds, or to alter equilibria, that can seriously damage sensitive chemical systems. Furthermore, practical problems may arise from small droplets sticking to a solid cover.

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An alternative is to diminish the solvent loss by controlling the environment in humidified chambers and by dispensing compensating solvent into the microvials via fine capillaries from above (Roeraade et al. (1996) Analytical Methods and Instrumentation. Special issue μ TAS'96 (1996), pp. 34-38). However, this technique can be ineffective over prolonged time periods and is subject to many practical problems associated with the restricted accessibility to the vials through the environmental chamber. Furthermore, since the solvent compensating capillaries block the space in close proximity to the microvials, accessing or detecting the material becomes increasingly more complex as the assay becomes larger.

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There is a need for microfluidic devices including a system for handling small volumetric amounts of liquid which avoids the above discussed drawbacks and allows for free access to the contained material, thus facilitating chemical manipulation of the liquid or the gaseous headspace environment and for monitoring of reaction products.

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A device having the features of claim 1 and a method having the features of claim 6 fulfill this need.

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30 BRIEF DESCRIPTION OF THE DRAWINGS

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Examples of embodiments of the present invention are illustrated in the accompanying figures, where:

5 Figure 1 is a schematic view of a first embodiment of a system in accordance with the present invention for containing small amounts of material in a droplet on top of the orifice of a microchannel;

10 5 Figure 2 is an enlarged view of the top of the capillary in Figure 1 illustrating a droplet;

15 Figure 3 illustrates the three different possible shapes of the liquid-gaseous interface;

10 20 Figure 4 illustrates a top of an embodiment of a microchannel in accordance with the present invention with a droplet and sample components immobilised on the microchannel rim;

25 15 Figure 5 illustrates a circular array of fabricated holes containing microdrops in accordance with other embodiment of the present invention;

30 30 Figure 6 is a sectional view of Figure 4 illustrating a solvent container in accordance with an embodiment of the present invention;

20 35 Figure 7 is a schematic view of a rectangular array in accordance with the present invention;

Figure 8 is a sectional view of the array of Figure 7.

25 40 DISCLOSURE OF THE INVENTION

45 30 The present invention provides a method for replacing solvent that is evaporating from a microvolume of solvent placed in an open microarea (MA) of a microfluidic device. The method has the characterising feature that replacement takes place continuously via a microchannel (2) that transports liquid to the microarea (MA) from a liquid reservoir (vessel). The method is particularly useful in the context of running reactions in the solvent present on the microarea (MA) in order to assay an analyte, for the synthesis of chemical compounds etc. The reactants used, including an

5 analyte and/or various reagents, may be soluble in the microvolume or immobilised
to a solid support in contact with the microvolume. The microarea (MA) may be the
10 orifice region of the microchannel and the microvolume in the form of a microdrop
(1), as shown in Figure 2. By continuously replacing the evaporated solvent via a
15 conduit (2) with solvent from a communicating vessel (3) the reactants present in
the microvolume are prevented from being desiccated. The sample is focused in the
microvolume as long as the evaporation rate of the solvent is higher than the
sample diffusion rate. It should be noted that the solvent compensating principle is
generally applicable to minute volumes, thus the liquid-gaseous interface may
20 10 appear in any of the different shapes illustrated in Figures 3 a)-c). In the case of
droplets shown figure 3 a), they can be formed by applying an overpressure to the
solvent supplying tubing. This causes the droplet size to be determined by the
diameter of the capillary orifice, the interfacial tension, the wettability of the capillary
25 15 material and the magnitude of the applied overpressure (which needs to be in
equilibrium with the interfacial pressure difference across the curved surface of the
droplet). The microarea (MA) can be located either, as illustrated in Figure 1, on top
of a single capillary (4), or as shown in Figures 5 – 8, as an array of microareas
30 20 carrying liquid in the form of drops (6) or liquid in the form of other physical
microappearances (9) (e.g. surfaces of the type shown in figures 3 b)-c)) formed on
top of an array of fabricated holes (7) each supplied from a common solvent
35 25 container (8). In the case of droplets, the overpressure needed can be created by
any means of pressure generation, e.g. from a hydrostatic head, a micropump or a
pressurised container.

40 25 The open geometry in this invention, with microareas carrying analyte- and/or
reagent-containing solvent in direct contact with the surrounding gaseous phase, is
favourable with respect to easy accessibility. For example, wet-chemical reactions
45 30 can easily be performed with sample components contained in the surface layers,
using reagents dispensed from external means directly to the microvolume of liquid
placed in the microarea, for instance from ink-jet dispensers or fine pipettes.
Furthermore, detection of analytes or reaction products can readily be made by
50 35 using optical detectors, such as CCD-cameras. Moreover, the equilibrium between
the solvent on the microarea and the surrounding gaseous phase could be

5 exploited for passive sampling of air-born constituents over prolonged time periods, thus enabling subsequent environmental analysis.

10 The solvents contemplated are often aqueous, i.e. consists of water, possibly mixed
15 with one or more water-miscible liquids, such as acetone, methanol, ethanol and isopropanol. This does not exclude the use of other solvents in the invention.

15 A second aspect of the invention relates to a microfluidic device comprising a
microchannel providing for liquid contact between an open microarea carrying a
20 microvolume of a solvent and a reservoir for the solvent, said reservoir and said
microchannel being adapted so that solvent evaporated from said microarea is
continuously replaced by solvent from the reservoir through said microchannel.
When in use the microvolume of solvent typically contains an analyte and/or one or
25 more reagents for assaying the analyte either directly or indirectly, for running
synthesis of a compound etc. By the term "indirectly" is contemplated that a feature
or an amount of a reaction product related to the analyte is assayed.

30 In order to avoid the risk of desiccation of the microareas over prolonged time
periods, the supplying solvent vessel should contain a solvent volume one, two
35 three or more orders of magnitude larger than the sum of all microvolumes
communicating with the reservoir.

35 The term "microvolume" means a volume that typically is at most around 10 μ l, such
as $\leq 1 \mu$ l. The lower end of the range extends down to the infinitesimal volume that
40 is present in the gaseous-liquid interface of the microvolume of the solvent.
Typically the microvolume is $\geq 10^{-15}$ l (femtolitre). It will be understood, however,
that the described principles may be applicable also to microvolumes being larger
45 than 10 μ l. By "microfluidic device" is meant a device that can handle
microvolumes, for example a volume that is less than 1 μ l, preferably between 1
50 and 10 nl, of reagents that may be introduced into the device.

50 A microarea may have different forms that vary from being an essentially flat form
via cup-formed areas to being walls of open chambers, the important matter being
that the area is able to carry the microvolume of liquid contemplated.

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Microchannels typically have the ability to act as capillaries. Normally their size in the dimension (i.e. height, width or length) in which they are smallest is less than 2000 μm , such as $\leq 500 \mu\text{m}$. Typically this dimension is $\geq 1 \mu\text{m}$. A microchannel 5 may be in form of a tube that may have a circular, a rectangular etc cross sectional area. They may also be "sheet"-like covering larger areas.

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The reagents included in or in contact with the microvolume of solvent vary depending on the reaction to be run. The reagents include catalysts, for instance, 10 an enzyme, compounds needed for the synthesis of nucleic acids, affinity reactants, etc. The term also includes biological systems, such as enzymatic systems and 20 whole cells. Affinity reactants typically form non-covalent complexes and may be illustrated by biotin, streptavidin, protein A, antibodies, lectins, hormone receptors, nucleic acids, peptides and polypeptides. Typical assays are immunoassays, 15 sequencing of nucleic acids and of peptides, hybridisation assays, detection of mutations, cell assays, etc.

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In one embodiment of the invention, one or more of the reagents used are immobilised in the microarea (MA). This alternative configuration is illustrated in 20 Figure 4, where reagents (11) are immobilised on the rim (5) of a microchannel, allowing washing steps to be performed by overflowing the microchannel. 35 Immobilisation may be achieved via covalent bonds, affinity bonds, physical adsorption etc. Typical affinity bonds are those formed by having strepavidin or a high affinity antibody bound to a solid support in the microarea (MA) and then 25 binding a desired reagent conjugated with biotin or with the hapten against which 40 the antibody is specific to the solid support bound strepavidin/high affinity antibody.

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The method for replacing solvents can be used in a method to prevent samples from becoming desiccated. One example of a method for achieving this comprises 30 the following steps:
providing a microarea for receiving a sample;
connecting the microarea (preferably via a microchannel) to a reservoir of 50 solvent;
applying the sample to the microarea;

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5 allowing solvent to evaporate from said microarea; and
10 continuously replacing said evaporated solvent with solvent from said reservoir.

15 In this example, it is preferable that the diffusion rate of the sample in the solvent
20 is less than the flow rate of solvent from the reservoir so that the sample does
not diffuse away from the microarea.

25 A second example of a method for preventing samples becoming desiccated
comprises the additional step of:
30 10 anchoring the sample to the microarea.

20 In this example, the flow rate of solvent from the reservoir may be less than the
diffusion rate of the sample in the solvent once the sample is firmly attached to the
microarea and is unable to diffuse away.

25 15 The sample can be applied to the microarea by dispensing from above, for
example by dropping into the microarea a drop of solvent containing the sample,
or from below, for example by injecting the sample into the microchannel
30 between the reservoir and the microarea and allowing the flow of solvent to bring
20 the sample to the micro area.

35 The microfluidic device according to the invention can suitably be fabricated in the
form of a circular (Figure 5 and 6) or rectangular array format (Figure 7 and 8),
although any other shape is also conceivable.

40 25 A circular format means that there are one or more microareas (chambers) that are
placed radially and in different directions from a centre. The distance from the
centre to individual microareas (chambers) may be equal or different. The reservoir
45 is preferably in the centre. The microchannels may be radially directed from the
30 centre and communicate with one or more microareas. The microchannels may
also be in the form of a common, flat-like microchannel or reservoir beneath the
microareas (chambers) and communicating upwardly via traditional microchannels.

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5 In rectangular formats there are microareas (chambers) that form a rectangular pattern. The microchannel arrangement may be in analogy with the circular format.

10 Microfluidic devices in the form of rotatable discs are known in the art. WO
15 97/21090 discloses a microanalytical / microsynthetic system for biological and
chemical analysis, comprising a rotatable microplatform, e.g. a disc, having inlet
ports, microchannels, detection chambers (microareas) and outlet ports through
20 which fluid may flow. Preferably, a circular array comprises a disc and a plurality of
microchannels (see Figure 5 and 6), each microchannel being radially dispersed
25 about the centre of the said rotatable disc. The rotatable disc is adapted for rotation
about its axis. Such adaptation may take the form of a hole at the axis of one or
both substrates which is capable of engaging a drive shaft. Other methods of
rotating the disc include clamping the disc and contacting the perimeter with a
30 moving surface, for example moving wheels, or placing the disc on a turntable and
spinning the turntable. Preferably the disc comprises a solvent inlet port located
35 towards the centre of the disc and connected to radially dispersed microchannels,
each microchannel having a sample reservoir located at the microchannel orifice
that is located outward from the centre of the disc.

40 20 The configuration of the microchannels in the rectangular or circular format may be
chosen to allow for application of a chemical compound, or a suspension of cells, to
35 the sample reservoir filled with fluid medium.

40 The microfluidic device may also comprise a separate microchannel system for
45 25 transporting one or more of the reactants needed to the microareas.

50 45 Suitably the circular or rectangular array format is a one- or two-piece construction
assembled together to provide a closed structure with openings at defined positions
to allow loading of the device with liquids and removal of waste liquids. In the
55 30 simplest form, see, for example, figures 6-7, the disc or wafer is produced as two
complementary parts (12), (13), one or each carrying channel structures which,
when affixed together, form a series of interconnected structures within the body of
a solid disc or wafer. The microchannels may be formed by micro-machining
methods in which the channels and chambers are micro-machined into the surface

5 of a disc or wafer, and a cover, for example a plastic film, is adhered to the surface so as to enclose the channels and chambers.

10 Suitable glass or polymeric materials can be additionally selectively modified by 5 chemical or physical means to alter the surface properties to confer a desired property, e.g. compatibility with cell growth, cell attachment and the attachment of biomolecules by covalent or non-covalent means.

15 Based on knowledge at the priority date, the variant given in figures 1 and 2 10 corresponds to the best mode in October 1998.

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Claims

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CLAIMS

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1. A microfluidic device comprising a microchannel (2) (4), providing for liquid contact between an open microarea (MA) or chamber carrying a microvolume (1) of a solvent and a reservoir (3); (8) for the solvent, said reservoir (3); (8) and said microchannel (2), (4) being adapted so that solvent evaporated from said microarea (MA) is continuously replaced by solvent from the reservoir (3); (8) through said microchannel (2), (4).

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2. The microfluidic device according to claim 1 wherein
 - a) said reservoir (3); (8) is positioned so as to create an overpressure in the solvent which is in equilibrium with the interfacial pressure difference across the curved surface of the droplet, or
 - b) said reservoir (3); (8) is connected to pump means that either facilitate replacement of solvent by pumping solvent or pressurising the reservoir (3); (8).

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3. The microfluidic device according to anyone of claims 1-2 comprising a plurality of microchannels (3); (8) and open chambers forming an array in the circular or rectangular format.

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4. The microfluidic device according to anyone of claims 1-3, wherein the microvolume contains one or more reactants that are soluble in the solvent or bound to a solid support in contact with the microvolume.
5. The microfluidic device according to claim 4 wherein at least one of said one or more reactants is an affinity reactant, for instance selected from nucleic acids, peptides, proteins.
6. A method for replacing solvents evaporating from a microvolume of solvent placed in an open microarea (MA) of a microfluidic device, characterised in that that replacement is continuously taking place via a microchannel (2), (4) that transports liquid to the microarea (MA) from a liquid reservoir (vessel) (3); (8).

5 7. The method of claim 6, characterised in that the microarea (MA), microchannel (2), (4) and reservoir are parts of the microfluid device defined in claims 1-5.

10 8. Method for replacing solvents for preventing samples from becoming desiccated characterised in that it comprises the following steps:
5 providing a microarea (MA) for receiving a sample;
 connecting the microarea (MA) to a reservoir (3); (8) of solvent;
 applying the sample to the microarea (MA);
 allowing solvent to evaporate from said microarea (MA); and
15 10 continuously replacing said evaporated solvent with solvent from said reservoir (3); (8).

20 9. Method in accordance with claim 8 characterised in that it comprises the additional step of:
25 15 anchoring the sample to the microarea (MA).

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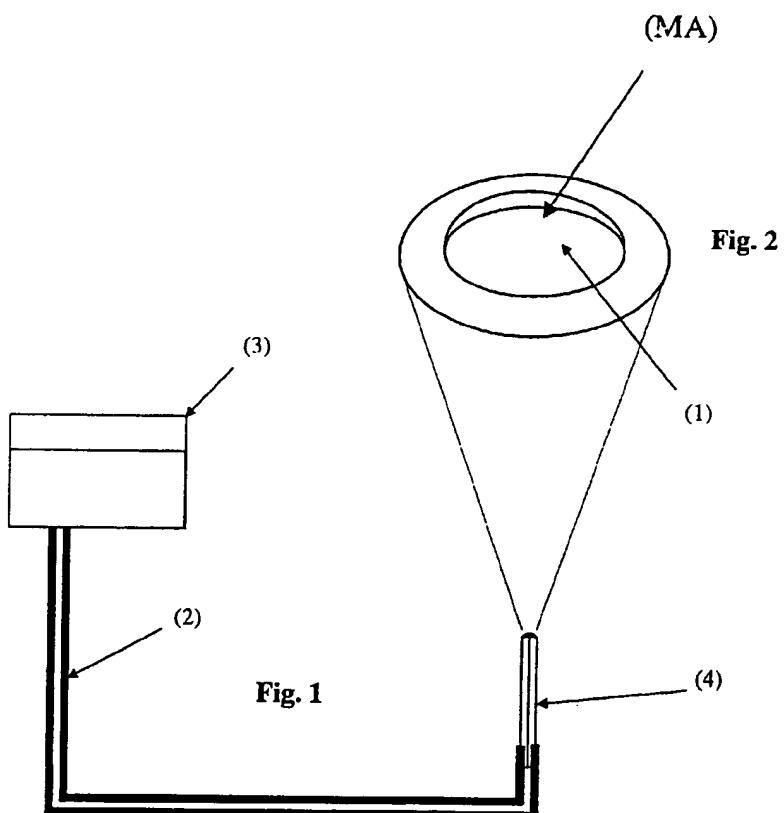


Fig. 3

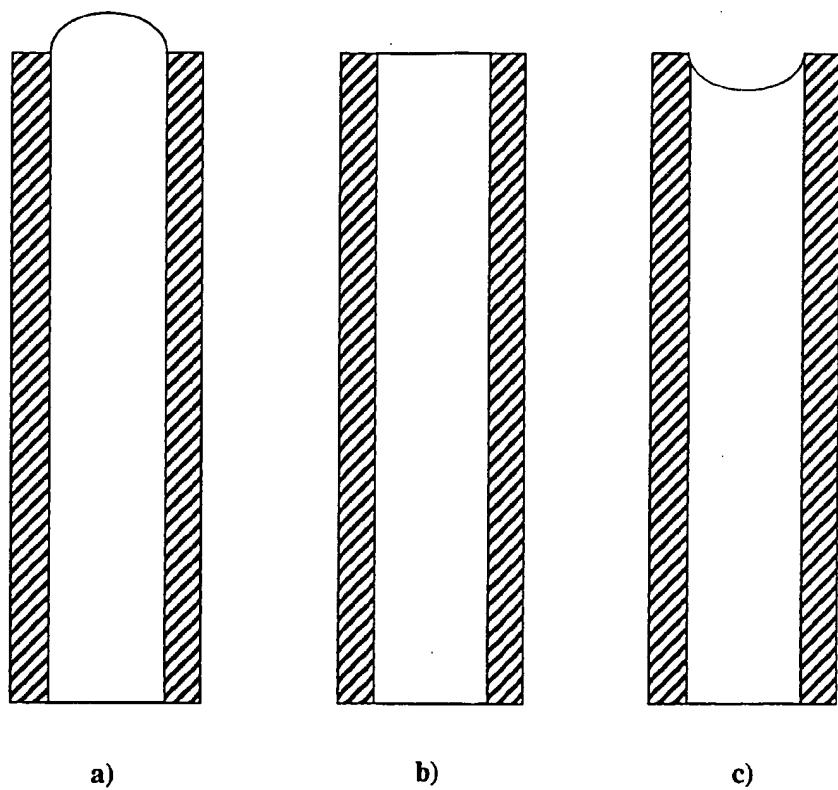


Fig. 4 (MA)

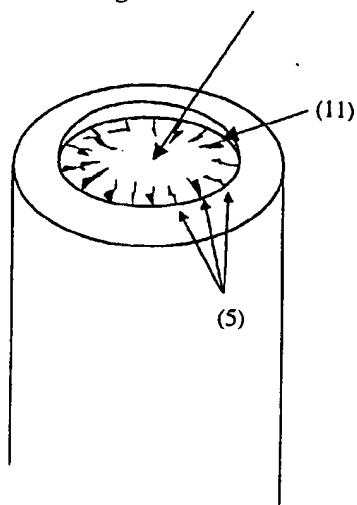


Fig. 5

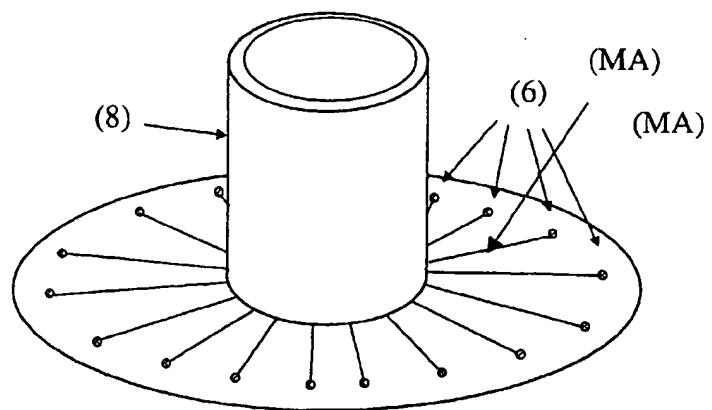


Fig. 6

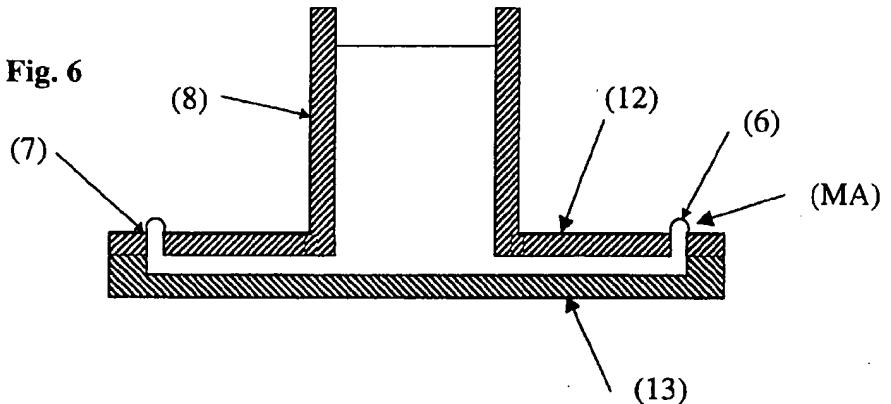
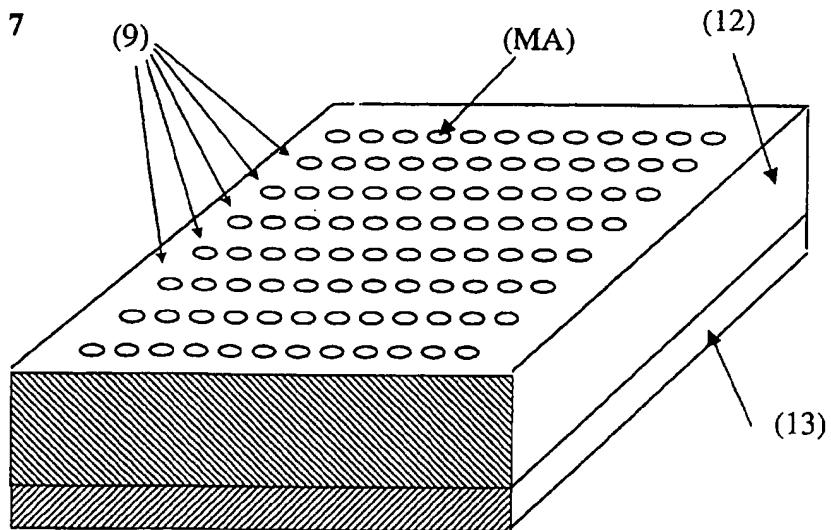
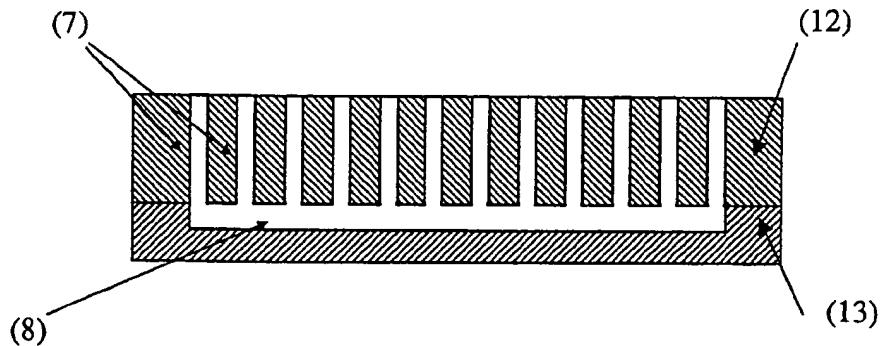


Fig. 7**Fig. 8**

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INTERNATIONAL SEARCH REPORTInternational application No.
PCT/SE 99/01958

A. CLASSIFICATION OF SUBJECT MATTER		
IPC7: B01L 3/00 According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED		
Minimum documentation searched (classification system followed by classification symbols)		
IPC7: B01L, G01N, C12Q, B01J Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched SE,DK,FI,NO classes as above		
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)		
WPI		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP 0305210 A2 (BIOTRACK, INC.), 1 March 1989 (01.03.89), figure 13 --	1-9
A	US 5783148 A (HUGH V. COTTINGHAM ET AL), 21 July 1998 (21.07.98), abstract --	1-9
A	WO 9855852 A1 (CALIPER TECHNOLOGIES CORP.), 10 December 1998 (10.12.98), figure 3 --	1-9
A	EP 0430248 A2 (MOCHIDA PHARMACEUTICAL CO., LTD.), 5 June 1991 (05.06.91), abstract -----	1-9
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INTERNATIONAL SEARCH REPORT
Information on patent family members

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International application No.
PCT/SE 99/01958

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